



PATENTS • DESIGNS  
The  
Patent  
Office  
• COPYRIGHT • TRADEMARKS •  
09/623 681

37C2

#5  
PCT/GB 00/03165

3726

INVESTOR IN PEOPLE  
REC'D 13 SEP 2000

WIPO PCT

The Patent Office  
Concept House  
Cardiff Road  
Newport  
South Wales  
NP10 8QQ

RECEIVED  
APR - 2 2001  
163700 MAIL ROOM

**PRIORITY  
DOCUMENT**

SUBMITTED OR TRANSMITTED IN  
COMPLIANCE WITH RULE 17.1(a) OR (b)

4

I, the undersigned, being an officer duly authorised in accordance with Section 74(1) and (4) of the Deregulation & Contracting Out Act 1994, to sign and issue certificates on behalf of the Comptroller-General, hereby certify that annexed hereto is a true copy of the documents as originally filed in connection with the patent application identified therein.

In accordance with the Patents (Companies Re-registration) Rules 1982, if a company named in this certificate and any accompanying documents has re-registered under the Companies Act 1980 with the same name as that with which it was registered immediately before re-registration save for the substitution as, or inclusion as, the last part of the name of the words "public limited company" or their equivalents in Welsh, references to the name of the company in this certificate and any accompanying documents shall be treated as references to the name with which it is so re-registered.

In accordance with the rules, the words "public limited company" may be replaced by p.l.c., plc, P.L.C. or PLC.

Re-registration under the Companies Act does not constitute a new legal entity but merely subjects the company to certain additional company law rules.

Signed

*R. Mahoney*

Dated

23rd August 2000

**THIS PAGE BLANK (USPTO)**

21 AUG 1999

## Request for grant of a patent

(See the notes on the back of this form. You can also get an explanatory leaflet from the Patent Office to help you fill in this form)

RECEIVED

21 AUG 1999

The Patent Office

Cardiff Road  
Newport  
Gwent NP9 1RH

1. Your reference XA1255

2. Patent application number  
(The Patent Office will fill in this part)

9919787.3

3. Full name, address and postcode of the or of each applicant (underline all surnames)

British Aerospace plc  
Warwick House, PO Box 87  
Farnborough Aerospace Centre  
FARNBOROUGH  
Hants GU14 6YUAdvanced Composites Group Ltd  
Composites House, Delves Road  
Heanor Gate Industrial Estate  
HEANOR  
Derbys DE75 7SJ6244933002  
Patents ADP number (if you know it)

If the applicant is a corporate body, give the country/state of its incorporation

51160092  
GB5710769001  
GB

4. Title of the invention

MANUFACTURE AND ASSEMBLY OF STRUCTURES

5. Name of your agent (if you have one)

GROUP IP DEPARTMENT

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

BRITISH AEROSPACE PLC  
LANCASTER HOUSE, PO BOX 87  
FARNBOROUGH AEROSPACE CENTRE  
FARNBOROUGH  
HANTS GU14 6YU

Patents ADP number (if you know it)

5116006

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (if you know it) the or each application number

Country      Priority application number  
(if you know it)      Date of filing  
(day / month / year)

N/A

7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application  
Date of filing  
(day / month / year)

N/A

8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if:

YES

- a) any applicant named in part 3 is not an inventor, or
- b) there is an inventor who is not named as an applicant, or
- c) any named applicant is a corporate body.

See note (d))

**Patents Form 1/77**

9. Enter the number of sheets for any of the following items you are filing with this form.  
Do not count copies of the same document

Continuation sheets of this form

Description	10
Claim(s)	3
Abstract	1
Drawing(s)	2 + 2

16

10. If you are also filing any of the following, state how many against each item.

Priority documents	0
Translations of priority documents	0
Statement of inventorship and right to grant of a patent ( <i>Patents Form 7/77</i> )	0
Request for preliminary examination and search ( <i>Patents Form 9/77</i> )	1
Request for substantive examination ( <i>Patents Form 10/77</i> )	0
Any other documents (please specify)	0

11.

I/We request the grant of a patent on the basis of this application.

Signature *Debra Lewis*  
D L LEWISDate  
20.8.99

12. Name and daytime telephone number of person to contact in the United Kingdom

DEBRA LEWIS  
01252 384633**Warning**

After an application for a patent has been filed, the Comptroller of the Patent Office will consider whether publication or communication of the invention should be prohibited or restricted under Section 22 of the Patents Act 1977. You will be informed if it is necessary to prohibit or restrict your invention in this way. Furthermore, if you live in the United Kingdom, Section 23 of the Patents Act 1977 stops you from applying for a patent abroad without first getting written permission from the Patent Office unless an application has been filed at least 6 weeks beforehand in the United Kingdom for a patent for the same invention and either no direction prohibiting publication or communication has been given, or any such direction has been revoked.

**Notes**

- If you need help to fill in this form or you have any questions, please contact the Patent Office on 0645 500505.
- Write your answers in capital letters using black ink or you may type them.
- If there is not enough space for all the relevant details on any part of this form, please continue on a separate sheet of paper and write "see continuation sheet" in the relevant part(s). Any continuation sheet should be attached to this form.
- If you have answered 'Yes' Patents Form 7/77 will need to be filed.
- Once you have filled in the form you must remember to sign and date it.
- For details of the fee and ways to pay please contact the Patent Office.

## MANUFACTURE AND ASSEMBLY OF STRUCTURES

The present invention relates to the manufacture and assembly of structures, particularly those structures having an outer layer or skin secured to or supported by a sub-structure and where the outer layer is required to conform within close tolerances to a predetermined profile.

It is often desirable to assemble such structures by first providing the sub-structure, or skeletal framework, and then attaching panels to the sub-structure to form the outer layer or skin.

This type of structure is common in the design and manufacture of aircraft where light-weight, high strength structures are required. In this industry it is additionally necessary to ensure that all the parts of the structure are manufactured within tight tolerances and furthermore that the parts fit together so that the assembled structure meets stringent accuracy requirements.

Any out of tolerance parts or inaccurately fitted parts will cause the adjacent parts being assembled to be fitted out of their required place in the structure, rendering the structure unacceptable. It is also important that adjacent parts of the outer skin do not have a step between them so that the constituent panels and skins of the outer layer of the aircraft structure are flush with one another. Failing to provide a substantially smooth aircraft outer layer can result in unwanted aerodynamic effects such as increased drag or turbulence.

To meet the strict tolerances required in aircraft construction for example the underlying sub-structure may be made from machinable aluminium or titanium. The sub-structure may then be machined as necessary to allow outer skins or panels to be fitted to it without adjacent skins or panels having a step between them. This method is not desirable as any machining errors may cause the whole sub-structure to fail a quality assurance inspection and be rejected with consequent cost and time penalties. Additionally underlying sub-structures are increasingly being made from lightweight composite materials such as carbon fibre reinforced plastics (CFRP) and these materials are not readily machinable.

A method of producing structures to high accuracy requirements is known, and can be used with sub-structures made of either metal or CFRP. In this method, the surfaces of sub-structure to which panels are to be attached are coated with a filled,

two component liquid adhesive material, with aluminium added to it. The liquid adhesive is cured on the sub-structure, and is then machined to a desired thickness before the panels or skins are fixed to the sub-structure. The cured adhesive may be machined to different thicknesses at different locations on the sub-structure so that, when the panels or skins are fixed to it there is substantially no step between adjacent panels or skins.

Whilst this method produces structures having profiles with acceptable accuracy, it has several disadvantages. Adhesive of this type is a viscous liquid which must be applied carefully to the sub-structure by hand using a spatula, so that it is distributed reasonably evenly with the desired thickness and without creating air bubbles in the adhesive. Too much adhesive will result in a longer wait for curing and more time spent in machining than necessary. Adhesive of this type is difficult to apply in desired quantities because of its viscosity and furthermore, there are health and safety implications associated with its use. Personnel must be trained to use such adhesive and must be careful when applying it to the sub-structure. Also special tooling must be manufactured, tailor made for each area to be panelled, to prevent the liquid adhesive from spreading to areas where it is not required, and to give guidance as to the thickness of the adhesive being applied. Because of the nature of this type of adhesive, the tooling must be coated with a release agent before use and cleaned thoroughly after use. Repeated exposure to this coating and cleaning process causes the tooling to deteriorate rapidly after a relatively low number of uses, resulting in time lost and expense in manufacturing and fitting replacement tooling.

The present invention seeks to provide a method of assembling structures having an outer layer supported by a sub-structure, and where the outer layer is required to conform, within close tolerances, to a predetermined profile, without the need to machine the sub-structure directly and avoiding the disadvantages of using a two-component liquid adhesive material.

The present invention further seeks to provide a machinable material for use as a shim between an outer layer and a supporting sub-structure.

The present invention additionally seeks to provide a method of making a material for use as a shim between an outer layer and a supporting sub-structure.

According to the present invention in one aspect thereof, there is provided a method of assembling a structure comprising at least the steps of

providing a sub-structure,

positioning shim material on at least part of the sub-structure,

curing the shim material disposed on the sub-structure,

machining the cured shim material to a desired thickness, and

assembling an outer layer with the sub-structure such that the shim material lies substantially between the outer layer and the sub-structure.

The shim material is preferably cured at low temperature, at below 80°C for example. The shim material is preferably in the form of a film or sheet.

Curing may be effected by exposure of the shim material to ultra violet light or radio frequency radiation.

The outer layer may comprise at least two parts, where each outer layer part is intended to be assembled with the sub-structure and each outer layer part is to be assembled substantially adjacent one or more other outer layer parts.

The shim material may be machined to different thicknesses at different locations on the sub-structure so that, when assembled to the sub-structure the outer layer parts together conform, within predetermined tolerances, to a predetermined profile. The thickness of each outer layer part is advantageously measured prior to machining the shim material.

The desired thickness of the shim material at a given location is dependent upon the thickness of the outer layer part which is to be assembled with the sub-structure at that location.

According to the present invention in another aspect thereof, there is provided a machinable shim material comprising at least a resin, a filler and a curing agent, wherein the shim material is capable of being cured at a temperature of below 80°C .

The resin may be any material capable of forming a stable matrix on curing. Suitable resins may be polyesters, urethanes, acrylics, epoxies, vinyl esters or phenolics, for example. Preferably the resin is an epoxy resin. The epoxy resin

advantageously comprises epoxy resin in both solid and liquid forms. Advantageously the epoxy resin is a Bisphenol A based epoxy resin.

The shim material is advantageously provided pre-formed, for example in the form of a film or a sheet. The film or sheet may be pre-cut so as to be suitable for direct use in particular applications, for example it may be supplied in pre-cut gasket form for use between parts having pre-determined matching dimensions.

Advantageously the shim material is thixotropic, giving minimal flow and slump but being capable of amalgamation at butted joints, at ambient temperatures of between 10°C and 35°C. The shim material preferably has viscosity such that it substantially does not flow during the heating process up to a cure temperature of 80°C. The shim material advantageously is capable of being stored for at least 6 months at -18°C in a stable state such that substantially no curing takes place.

The filler may be an inert material such as talc, calcium carbonate, aluminium silicate, dolomite, alumina trihydrate, wollastonite or glass fibres. In addition the filler may comprise light weight glass microballoons, cenospheres, phenolic microballoons, perlite, hollow ceramic spheres or other plastic spheres. In addition the filler may comprise metallic substances such as aluminium powder, air release agents such as silanes or surfactants, and adhesion promoters such as organosilanes or titanates. The filler is advantageously chosen to be capable of aiding a subsequent machining process.

The shim material preferably comprises a thixotropic agent such as, for example, silicon dioxide, hydrogenated castor oil, sepiolite, bentonite clays, attapulgite clays or diatomaceous earths.

Advantageously the curing agent is a substituted imidazole. Alternatively the curing agent may be an amine, a boron trifluoride complex or a dicyandiamide with a substituted urea accelerator. Preferably the curing agent is 1-methyl imidazole. The curing agent is preferably present at a level of 2 to 10 parts per hundred by weight.

The shim material may be used in the method of producing structures described above.

According to the present invention in a further aspect thereof, there is provided a method of making a shim material comprising at least the steps of

- 5 -

heating a resin to at least 80°C,  
adding a filler to the resin,  
cooling the resin to below 60°C,  
adding a curing agent to the resin, and  
forming the resin into a sheet of desired thickness.

The method may be used to make the preformed machinable shim material described above.

Where the resin is an epoxy resin it is preferably heated to at least 100°C.

Where a thixotropic agent is to be included this may be added with the filler to the resin before it is cooled. Where silicon dioxide is used as the thixotropic agent it is preferably in the form of fumed silica.

Aluminium powder may be added to the epoxy resin following heating and before cooling.

An air release agent may be added to the epoxy resin following heating and before cooling.

Reodourant may be added to the epoxy resin following cooling if desired.

Adhesion improving agents, for example organosilanes, titanates or modified rubbers, may be added to enhance the adhesive quality of the shim material if desired.

The sheet of shim material may be stored in the freezer until it is required for use.

Embodiments of the invention will now be described, by way of example only, and with reference to the following drawings and examples.

Figure 1 shows an exploded isometric sketch of a sub-structure and parts for assembly into an aircraft structure,

Figure 2 shows a section through a prior art assembly of the sub-structure and parts shown in Figure 1,

Figure 3 shows a section through an assembly of the sub-structure and parts shown in Figure 1 utilising an embodiment of the present invention,

Example 1 describes a material suitable for use as shim material, for example in the assembly shown in Figure 3, and

Example 2 describes a method of producing the material of Example 1.

Referring now to the drawings in which the same features are denoted by common reference numerals:

Figure 1 shows a structure 1 of part of an aircraft comprising a sub-structure 2 and several outer layer parts 4, 6, 8, 10, 12 to be fitted to the sub-structure 2. The parts and the sub-structure may be formed from metal or carbon fibre composite, as desired. The parts 4, 6, 8, 10, 12 and sub-structure 2 are all manufactured and assembled with adherence to strict tolerance limits.

Figure 2 shows part of the structure 1 comprising parts 4 and 8 which form part of the outer layer of an aircraft. Parts 4 and 8 may be outer skins or detachable panels. Using a prior art method of assembly the parts 4, 8 are fixed to sub-structure 2 with a gap 14 between them. Parts 4, 8 are wet assembled to the substructure 2 using PRC (polysulphide rubber compound) to prevent liquid ingress, the PRC seeping into the gap 14 during assembly. The PRC is applied to the sub-structure before the parts 4 and 8 are assembled to it.

Due to the cumulative dimensional tolerances of the sub-structure and the parts resulting from their individual manufacture, there is a step D between adjacent parts 4 and 8, which results in unwanted aerodynamic effects which can affect the performance of the aircraft. For aircraft applications it is typically preferable to have a step D of less than 0.5mm between adjacent parts.

Figure 3 shows part of the same structure 1 comprising parts 4 and 8 which form part of the outer layer of an aircraft structure. As before the parts 4 and 8 may be outer skins or detachable panels and in this example are designed to be of different thicknesses, both panels being within their respective tolerances. The sub-structure 2 is designed to be machined to a profile which is designed to accommodate the different design thicknesses of parts 4 and 8 such that, when assembled, parts 4 and 8 should be substantially flush with one another, so that steps such as D shown in Figure 2 are no greater than 0.5mm. The step E represents the difference in thickness between parts 4 and 8. However, due to variation of the substructure profile, and the cumulative dimensional tolerances of the

substructure and the parts 4, 8 a step D would inevitably occur if the prior art method of assembly described above were to be used. However to counter step D, a layer of shim material 20 is introduced between sub-structure 2 and the parts 4 and 8. The shim material is machined to allow parts 4, 8 to be substantially flush when assembled on the substructure, reducing step D to less than 0.5mm.

Using our new method of assembly the layer of shim material 20 of 2mm thickness is stuck to the sub-structure 2. The shim material 20 is adapted to have a tackiness which allows it to stick to surfaces on which it is positioned, even if the surfaces are vertical or on the underside of the sub-structure. The shim material may be cured to a machinable condition at 21°C under atmospheric pressure in 12 hours. Alternatively the shim material may be cured to a machinable condition in 1 hour at 65°C. The heating may be achieved by a hot air gun, using hot rollers, an oven, bank of heaters, a gas thermocatalytic heater or other conventional heating methods.

The shim material substantially does not flow when heated, and so tools for providing boundaries are not required for this method of assembly.

After the shim material 20 is cured, it is machined using conventional machine tools by a desired amount. To determine the desired amount, the actual thicknesses of parts 4 and 8 are first determined and the location on the shim material 20 on which the parts will be positioned is identified. In this example part 4 is to be positioned at location 18 and part 8 at location 16. The shim material 20 is then machined by differing amounts at locations 18 and 16 respectively, so that the actual profile of the assembly is within tolerance of the datum profile of the assembly. It should be noted that any variations in the surface of the sub-structure 2 or thickness of parts 4 or 8, which might contribute to a step between attached parts 4 and 8, are substantially eliminated by the application of the shim material 20 and its subsequent precision machining.

The shim material 20 is kept in a freezer to prevent curing until it is required for use.

The shim material does not chemically react or otherwise interfere with sealant, such as cured PRC or a cured two part adhesive material, used as a filler to fill cracks and voids in the substructure.

- 8 -

The shim material is self-amalgamating, i.e. when strips of shim material laid side by side adjacent each other they coalesce. Self amalgamation is temperature and time dependent and preferably occurs within 1 hour at 65°C. The flow characteristics of the shim material may be altered by varying the relative amounts of ingredients.

A material suitable for use as a shimming material, and which may be used in the method described with reference to Figure 3 is a thixotropic filled resin material with a specifically controlled tack which allows strong adherence to vertical or inverted surfaces and facilitates full wetting of the substructure without air entrainment, the filled resin material incorporating a curing agent. An example of such material is as follows

EXAMPLE 1	
Ingredient	Parts by Weight
Bisphenol A based epoxy resin in liquid form	88
Bisphenol A based epoxy resin in solid form	12
Talc filler	40
Hollow glass microspheres	25
Fumed silica (thixotropic agent)	6
Aluminium powder	5
Air release agent	0.5
1 – Methyl imidazole (curing agent)	6
Reodourant (if desired)	0.1
Adhesion Improving Agent (Organosilane)	0.1

An example of a method for producing the material described in Example 1 will now be described by way of example only.

#### Example 2

Pre-heat the Bisphenol A epoxy resins to a temperature at which the viscosity is sufficiently lowered for filler to be incorporated. This temperature will be dependent

upon the nature of the epoxy resins and the power rating of the mixing equipment, and will be typically approximately 120-160°C.

Add in the talc filler, heat in a vacuum to degas the mixture and stir.

Add in the glass spheres, degas the mixture and stir.

Add in the air release agent, the adhesion improving agent and the aluminium powder and degas the mixture.

Add in the fumed silica.

Cool the mixture to below 60°C (preferably to 48°C) and degas.

Add the curing agent.

Discharge the mixture, a rollable dough-like substance, on to film forming equipment such as a reverse roll coater, knife over roller coater, extruder or conveyor press.

Form film of desired thickness, incorporating a supporting scrim material if required, and keep the shim material at refrigerated temperatures to prevent curing, until the shim material is required for use.

The shim material may be supplied in suitable lengths of material of selectable starting thickness, or it may be supplied in the form of pre-cut "gaskets" designed to fit the surfaces of the sub-structure to accommodate panels of various shapes. It should be curable at 65° and have a useable life out of refrigeration of at least 12 hours. The scrim material improves the fatigue life of the shim material, and strengthens the shim material. The scrim material ideally weighs 10-50 g/m<sup>2</sup> and is incorporated into one side of the film during manufacture of the shim material. The scrim material is preferably brightly coloured so that an operator can see which side of the film incorporates the scrim material and use it as a 'witness' mark during machining of the shim. During use, it is advantageous to position the film such that the supporting scrim material is adjacent the sub-structure and so does not get machined away following curing of the shim material.

It will now be apparent to one skilled in the art that characteristics of the shim material may be altered to suit different applications by changing the relative amounts of ingredients. For example, to alter flow characteristics or adhesive characteristics the ratio of filler to resin may be varied. As another example, the density of the shim

- 10 -

material may be altered by varying the amounts of different filler materials relativ each other.

Suitable ranges for various ingredients of the shim material are indicated below.

INGREDIENT	PERCENTAGE (%) BY WEIGHT
Bisphenol A based epoxy resin in liquid form	20-80%
Bisphenol A based epoxy resin in solid form	0-50%
Talc filler	0-70%
Hollow glass microspheres	0-70%
Fumed silica	0-10%
Aluminium powder	0-50%
Air release agent	0-3%
1-Methyl imidazole	2-10%
Reodourant	0-1%
Adhesion Improving Agent	0-2%

**CLAIMS**

1. A method of assembling a structure comprising at least the steps of:
  - providing a sub-structure,
  - positioning shim material on at least part of the sub-structure,
  - curing the shim material disposed on the sub-structure,
  - machining the cured shim material to a desired thickness, and
  - assembling an outer layer with the sub-structure such that the shim material lies substantially between the outer layer and the sub-structure.
2. A method as claimed in claim 1 wherein the shim material is cured at below 80°C.
3. A method as claimed in any one preceding claim wherein the curing is effected by exposure of the shim material to ultra violet light.
4. A method as claimed in claim 1 or claim 2 wherein the curing is effected by exposure of the shim material to radio frequency radiation.
5. A method as claimed in any one preceding claim wherein the outer layer comprises at least two parts and the thickness of each outer layer part is measured prior to machining the shim material.
6. A method as claimed in claim 5 wherein the shim material is machined to different thicknesses at different locations on the sub-structure so that, when assembled to the sub-structure, the outer layer parts together conform, within pre-determined tolerances, to a pre-determined profile.
7. A method of assembling a structure substantially as hereinbefore defined and with reference to the accompanying drawings and description.
8. A machinable shim material comprising at least a resin, a filler and a curing agent, wherein the shim material is capable of being cured at a temperature of below 80°C.
9. A shim material as claimed in claim 8 wherein the resin is an epoxy resin.
10. A shim material as claimed in claim 9 wherein the epoxy resin is a Bisphenol A based epoxy resin.

- 12 -

11. A shim material as claimed in any one of claims 8 to 10 wherein the shim material is in the form of a film or sheet.
12. A shim material as claimed in any one of claims 8 to 11 wherein the shim material is thixotropic.
13. A shim material as claimed in any one of claims 8 to 12 wherein the shim material has viscosity such that it substantially does not flow during the heating process up to a cure temperature of 80°C .
14. A shim material as claimed in any one of claims 8 to 13 wherein the shim material is capable of being stored for at least 6 months at -18°C in a stable state such that substantially no curing takes place.
15. A shim material as claimed in any one of claims 8 to 14 wherein the shim material comprises an air release agent.
16. A shim material as claimed in any one of claims 8 to 15 wherein the shim material comprises an adhesion promoter.
17. A shim material as claimed in any one of claims 8 to 16 wherein the curing agent is a substituted imidazole.
18. A shim material as claimed in any one of claims 8 to 17 wherein a curing agent is 1 – methyl imidazole.
19. A shim material as claimed in any one of claims 8 to 18 wherein the curing agent is present at a level of two to ten parts per hundred by weight.
20. A shim material substantially as hereinbefore defined and with reference to the accompanying drawings and description.
21. A method of making a shim material comprising at least the steps of:
  - heating a resin to at least 80°C,
  - adding a filler to the resin,
  - cooling the resin to below 60°C,
  - adding a curing agent to the resin, and
  - forming the resin into a sheet of desired thickness.

- 13 -

22. A method of making a shim material substantially as hereinbefore defined and with reference to the accompanying drawings and description.

## ABSTRACT

The invention relates to a shim material for use in assembling structures and a method of producing the shim material.

The shim material is machinable and comprises at least a resin, a filler and a curing agent, and is capable of being cured at a temperature of below 80°C.

The shim material is produced by heating the resin to at least 80°C, adding the filler to the resin, cooling the resin to below 60°C, adding a curing agent to the resin, and then forming the resin into a sheet of desired thickness.

Structures, such as airframe structures may be assembled using the shim material by positioning shim material on at least part of a sub-structure, curing the shim material in situ, machining the cured shim material to a desired thickness, and then assembling outer layer parts to the sub-structure with the shim material between structure and parts.

1/2

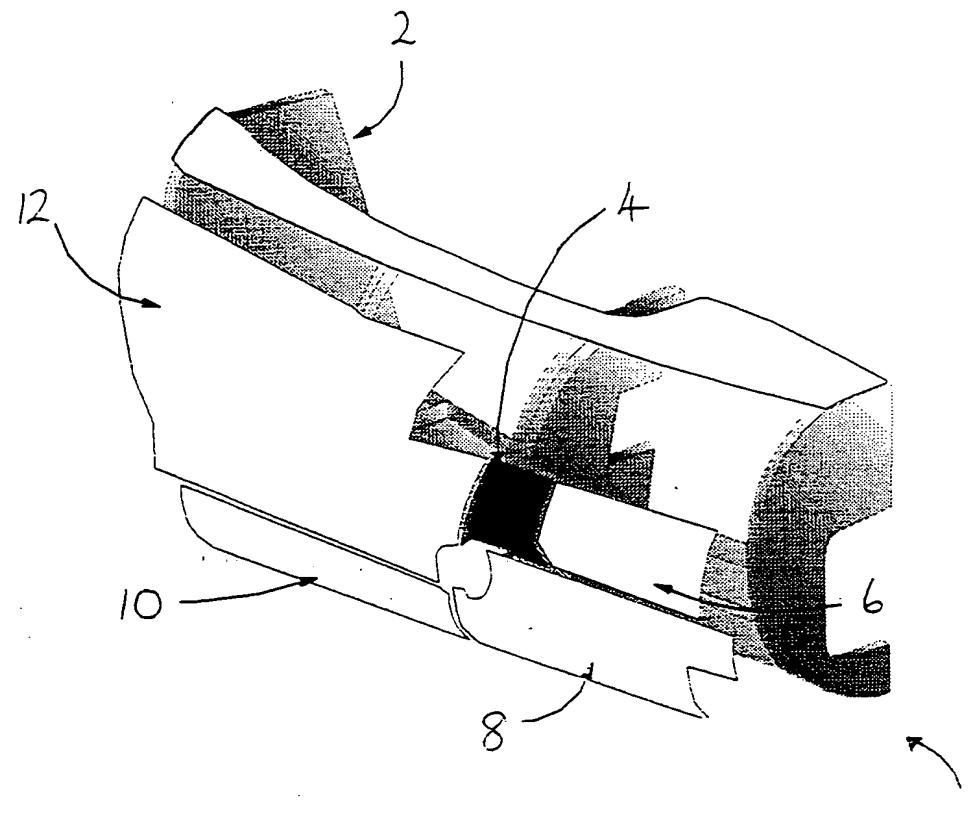


Figure 1

**THIS PAGE BLANK (uspto)**

2/2

PRIOR ART

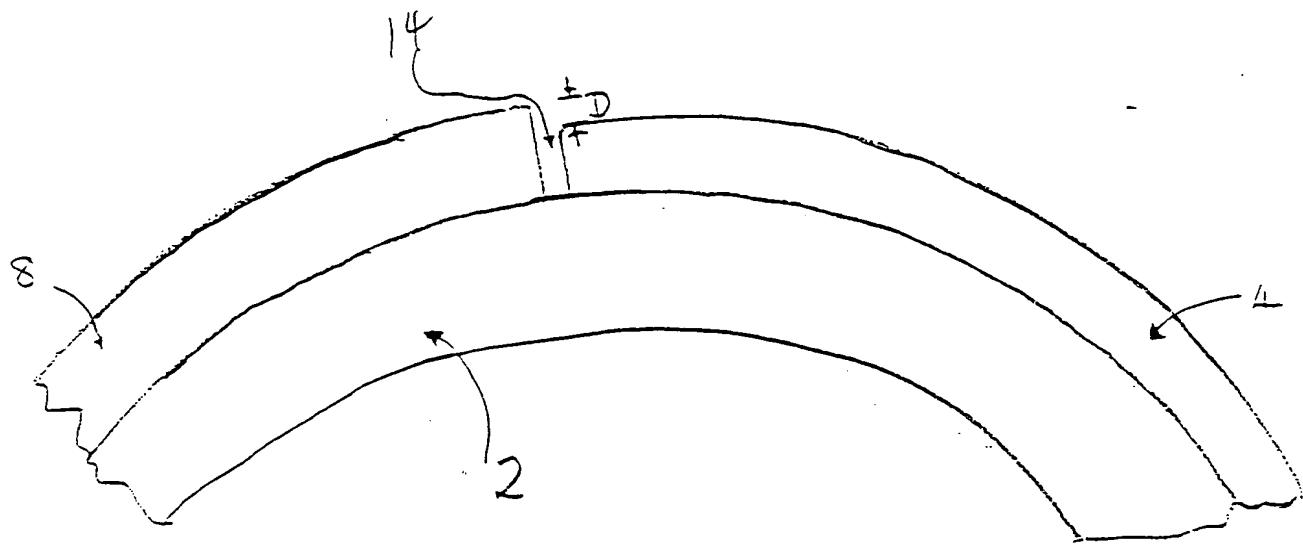


Figure 2

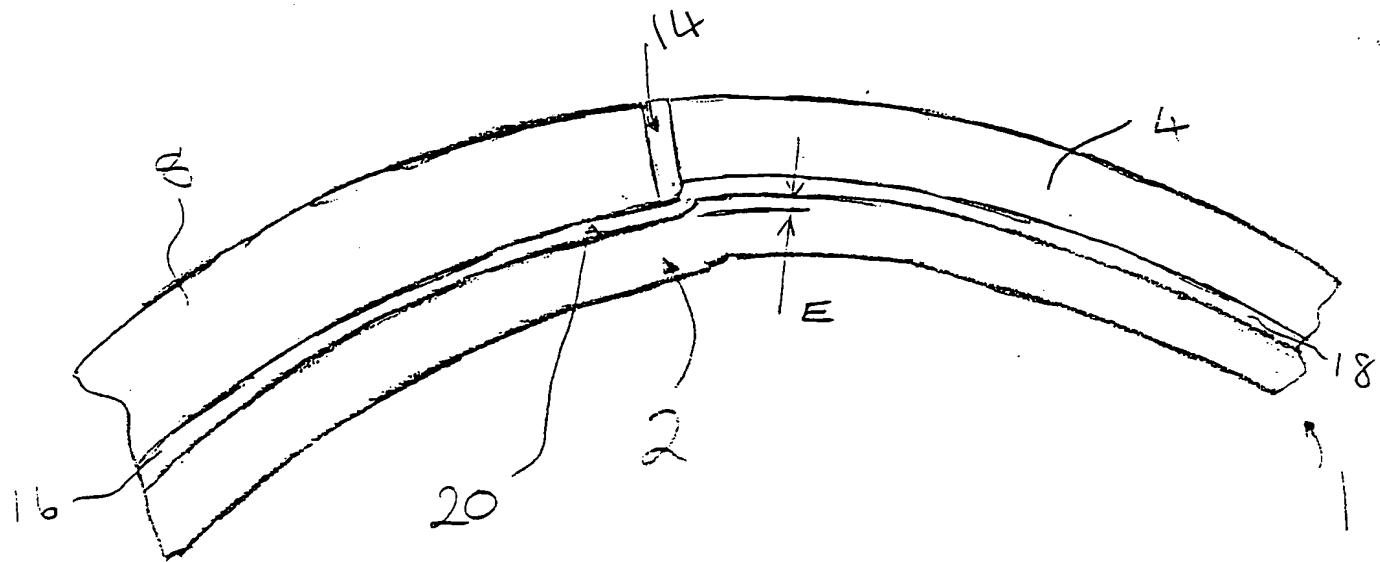


Figure 3

PCT/GB 05/03183

2377 filed 16-8-2000

BAE Systems plc

**THIS PAGE BLANK (USPTO)**